

Spinal meningeal melanocytoma with direct bone metastasis: A case report and literature review

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ABSTRACT

Meningeal melanocytoma is a rare benign tumor of the central nervous system. An intradural extramedullary tumor at C2 was accidentally discovered in a 58-year-old woman six years ago. The lesion was well-circumscribed mass on computed tomography and magnetic resonance imaging. We predicted the tumor to be a meningioma. We performed tumor resection because its gradual growth compressed the spinal cord. Intraoperatively, the tumor was dark brown and the nearby dura matter and the lamina had a pigmented lesion suggesting direct invasion. After gross total removal, the tumor was diagnosed as meningeal melanocytoma by pathological examination. Meningeal melanocytoma with direct bone metastasis are rare. We present this case with reference to previous literature.

Keywords: Direct bone metastasis, Melanoma, Radiological features, Spinal meningeal melanocytoma, Spinal tumor

How to cite this article

Inomo T, Aoyama M, Maejima R, Hara M. Spinal meningeal melanocytoma with direct bone metastasis: A case report and literature review. Int J Case Rep Images 2023;14(2):80–84.

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Received: 07 August 2023

Accepted: 15 September 2023

Published: 03 October 2023

Article ID: 101415Z01TI2023

doi: 10.5348/101415Z01TI2023CR

INTRODUCTION

Melanocytomas comprise <0.1% of all tumors in the central nervous system (CNS) [1]; meningeal melanocytoma is a rare benign tumor derived from leptomeningeal melanocytes [2]. Almost all of these tumors are found at the posterior fossa or upper spinal level [3, 4]. The most common image appearance is as a well-defined intradural extramedullary mass; other differential diagnoses are meningioma or schwannoma [3, 5]. Typical magnetic resonance imaging (MRI) findings are hyperintense on T1-weighted images (T1WI) and hypointense on T2-weighted images (T2WI), with unclear contrast enhancement [3]. In pathology, melanoma should be excluded. Relatively good prognosis can be expected after total resection. However, some cases report an aggressive course of rapid growth, recurrence, and transformation to malignant melanoma [6, 7]. Accordingly, careful observation is essential. We report a rare case of meningeal melanocytoma with direct bone invasion at the C2 level that was completely removed by surgery.

CASE REPORT

A 52-year-old woman with no previous medical history was accidentally diagnosed with a cervical spinal tumor at C2 level when injured in a traffic accident (Figure 1A). Since then, she had been complaining of numbness of the left limbs but there was no clear association with the lesion. The tumor kept gradually growing and the spinal cord became compressed over a 6-year follow-up, without symptom exacerbation. She planned to undergo tumor removal in August 2022. Preoperatively, the mass was well-circumscribed, and no calcification was observed by

computed tomography (CT). A cervical spinal CT revealed that it was intradural extramedullary, hyperintense on T1WI, hypointense on T2WI, with unclear contrast enhancement (Figure 1B–H). There was no lesion on head MRI.

A posterolateral approach was performed to resect the tumor. We approached the dura matter by C2 partial laminectomy and performed an incision. Intraoperatively, we found that the tumor was dark brown, and the lamina of the C2, dura, and pia nearby were partially pigmented. We removed the tumor piece by piece due to its softness and adhesion to dura, until achieving gross total resection (Figure 2A–E). After resection, the dural attachment was coagulated and the dura was sutured water-tight.

In pathological examination, proliferation of tumor cells with high melanin deposition was found in the tumor and in the pigmented lesion of the vertebral arch and dura (Figure 3A). There was slight cellular atypism but no atypical mitotic figure (0/10 high power field) but no clear necrosis. Immunohistochemistry analysis showed positive

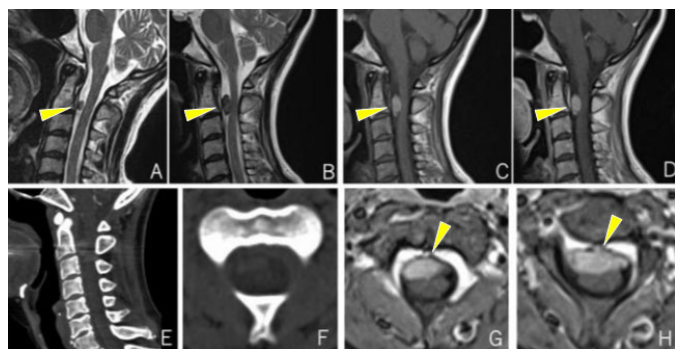


Figure 1: Preoperative cervical MRI and CT. (A) Sagittal T2-weighted image showing an extradural mass at C2 level at first diagnosis (yellow arrow head). (B) Sagittal T2-weighted image after 6 years of follow-up. Tumor gradually growing and compressing the spinal cord (yellow arrow head). (C) Sagittal T1-weighted image (yellow arrow head). (D) Sagittal T1-contrast-enhanced image (yellow arrow head). (E) Sagittal contrast-enhanced CT. (F) Axial contrast-enhanced CT. (G, H) Axial T1-weighted contrast-enhanced image (yellow arrow head).

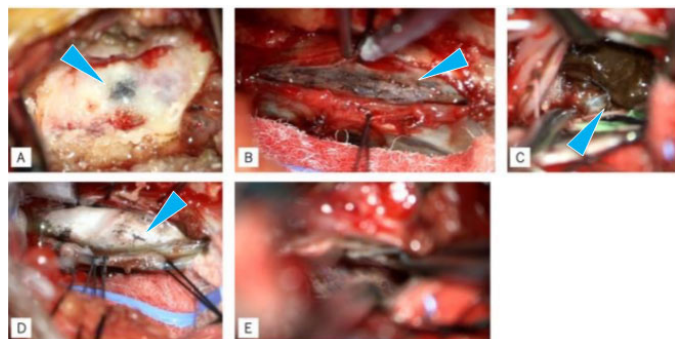


Figure 2: Intraoperative microscopic view. (A) A bone melanin lesion in the C2 vertebral arch (blue arrow head). (B) Pigmented arachnoid (blue arrow head). (C) A black colored tumor, soft and adhered to the dura (blue arrow head). (D) The pia nearby the tumor also pigmented (blue arrow head). (E) Complete gross total resection.

results for SRY-related high-mobility-group/HMG box 10 (SOX-10), S-100 protein, and human melanoma black-45 (HMB-45) (Figure 3B–E). The Ki-67 index was <2% (Figure 3F). The tumor was diagnosed as meningeal melanocytoma as there were insufficient findings to consider it as malignant melanoma. The patient is followed up regularly but not receiving radiation or chemotherapy. An MRI at three months after surgery showed no evidence of tumor recurrence (Figure 4A and B).

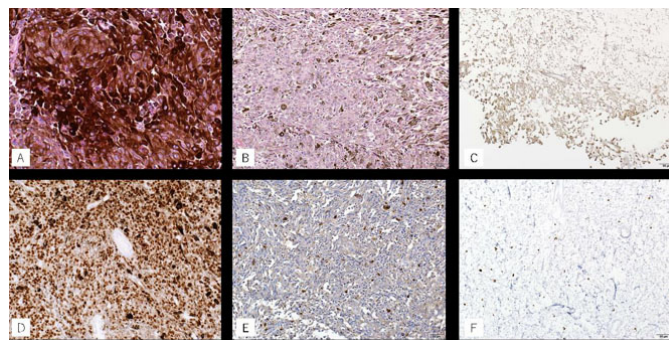


Figure 3: Histopathological and immunohistochemical findings. (A) Hematoxylin and eosin (H&E) staining (×400) showing high melanin pigmentation. (B) H&E staining after demelanization (×200). (C) HMB-45(+) staining of tumor cells (×200). (D) SOX-10(+) staining of tumor cells (×200). (E) S-100(+) staining of tumor cells. (F) Ki-67 staining showing a proliferation rate <2% (×200). HMB-45: a monoclonal antibody that reacts against an antigen present in melanocytic tumors such as melanomas. Ki-67: a marker to determine the growth fraction of a given cell population. SOX-10: a nuclear transcription factor that participates in neural crest development and in the differentiation of cells of melanocytes and Schwann cells. S-100: a protein presented in cells derived from the neural crest such as Schwann cells and melanocytes.

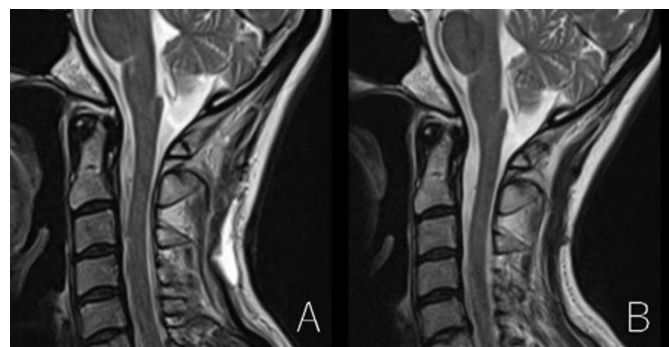


Figure 4: Follow-up MRI. (A) Sagittal T2-weighted image after surgery. (B) Sagittal T2-weighted image at three months after surgery.

DISCUSSION

In 1972, Limas and Tio first reported a melanocytic tumor in the craniocervical junction [8]. According to the 2016 World Health Organization classification of CNS tumors, primary melanocytic tumors of the CNS are classified as: (1) meningeal melanocytosis (Grade 0),

(2) meningeal melanocytoma (Grade 1), (3) meningeal melanoma (Grade 3), and (4) meningeal melanomatosis (Grade 3). Surgical removal is recommended as primary treatment [9]. Gross total removal is not difficult for intradural extramedullary tumors. The radiographic features and treatment of primary CNS melanocytoma from 10 case reports are summarized in Table 1. Most cases (9/10) achieved gross total resection or subtotal/near-total resection. In one case of partial resection, the patient was treated by radiation, immunotherapy, and second surgery, but he finally expired three years after diagnosis [15]. Two cases suffered recurrence and malignant transformation despite gross total resection and underwent chemotherapy. However, they showed resistance to the treatment [6, 16]. To the best of our knowledge, there are few reports of a good course with radiation and chemotherapy. We highly recommend total resection as first treatment of meningeal melanocytoma before tumor growth to prevent complications. Radiation and chemotherapy should be recognized as adjunctive treatment when total resection is not achieved or in case of tumor recurrence. Recently, immunotherapy with ipilimumab (anti-cytotoxic T1-Lymphocyte-associated protein drug) or nivolumab (anti-programmed cell

death protein-1 drug) showed good prognosis in melanoma [18]. The suggested effectiveness of those treatments for meningeal melanocytoma or melanoma warrants further research [16, 18]. Histopathologic and immunohistochemical examination is essential to diagnose meningeal melanocytoma. Immunohistochemical staining indicates that melanocytoma is positive for S-100 and HMB-45 but negative for epithelial membrane antigen [3, 19, 20]. In addition, the Ki-67/MIB-1 index is generally low (<5%), in contrast with the high score found in melanoma [2, 3].

In our case, the tumor was well-circumscribed, hyperintense, on T1WI and hypointense on T2WI, with unclear contrast enhancement. Its slow growth as well as the histopathologic and immunohistochemical findings of low Ki-67 index and no abnormal mitosis/necrosis suggested meningeal melanocytoma. In contrast, the intraoperative findings of direct bone metastasis cannot exclude melanoma. Previous research indicated a transformation of melanocytoma into melanoma [6, 7, 16], so we should consider radiotherapy depending on the disease course. Despite some reports of dural invasion in meningeal melanocytoma, we found no report of direct bone invasion; thus, the present case is rare.

Table 1: Cases of primary meningeal melanocytoma located in spinal cord in the literature

Author	Age/ Sex	Location	Clinical manifestation	T1	T2	Gd	Operation	Radiation (R) Chemotherapy (C) Immunotherapy (I)	Ki-67	Follow- up after surgery (month)	Recurrence	Outcome
Ok Hwa Kim [10]	37/M	C5-6	Posterior neck pain and right upper limb pain	Hyper	Iso	Iso	Subtotal resection	R	<1%	2	2 weeks after surgery	Death
Ahmed M. Salah El-Din [11]	5/M	T11-L4	Back pain	Hyper	Iso	Homogeneous enhancement	Subtotal resection	–	<1%	1	–	Alive
Daniele Armocida [12]	60/M	T10	Back pain and right hemiparesis	Hyper	Iso	Homogeneous enhancement	Subtotal resection	–	2%	1	–	Alive
Isamu Miura [13]	40/M	C2-3	Numbness of right upper limb and left lower limb	Hyper	Iso	Iso	Total resection	–	<1%	6	–	Alive
Makoto Tateyama [14]	27/M	C1-2	Posterior neck pain	Hyper	Iso	–	Near-total resection	R	–	36	1 year after surgery	Alive
Virginie Hean [15]	70/M	C7-T1	Sensory and motor disfunction of left lower limb	Hyper	Iso	Homogeneous enhancement	Partial resection	R/I	<5%	36	Growth	Death
Salman T. Shaikh [16]	36/F	L3-4 Intervertebral foramen	Back pain	Hyper	Iso	Homogeneous enhancement	Total resection	R/I	3–4%	48	6 months after surgery and malignant transformation	Death
Koichi Uramaru [17]	44/M	FM-C1	Headache and nausea	Hyper	Iso	Iso	Total resection	–	4%	24	–	Alive
Shuang-lin Deng [6]	19/F	C1-2	Headache, limbs weakness, and dyspnea	Hyper	Iso~ Hyper	Homogeneous enhancement	Total resection	R/C/I	1–2%	Unclear	6 months after surgery and malignant transformation	Death
Our Case	58/F	C1-2	Numbness of left upper limb	Hyper	Iso	Iso	Total resection	–	<2%	3	–	Alive

CONCLUSION

We report a rare case of meningeal melanocytoma in the upper cervical spine. Before surgery, we considered the tumor as a meningioma. It was difficult to identify as meningeal melanocytoma from images and clinical symptoms, as intraoperative findings, pathological examination, and immunostaining are essential to diagnose it. We recommend complete excision at an early stage as first choice treatment to avoid complications and transformation to malignant melanoma. In the present case, the gradual growth and pathological appearance indicates benign and good prognosis from surgical total resection. However, the direct bone invasion makes local recurrence possible, so regular follow-up is mandatory.

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Author Contributions

Toshiaki Inomo – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Masahiro Aoyama – Conception of the work, Design of the work, Drafting the work, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Ryuya Maejima – Conception of the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Masahito Hara – Acquisition of data, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related

to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Guarantor of Submission

The corresponding author is the guarantor of submission.

Source of Support

None.

Consent Statement

Written informed consent was obtained from the patient for publication of this article.

Conflict of Interest

Authors declare no conflict of interest.

Data Availability

All relevant data are within the paper and its Supporting Information files.

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